PTC/SB/08x (08-03)
Approved for use through 97/31/2006 ONE 0651-0001
U.S. Patent and Trademark Office, U.S. DEPARTMENT OF COMMERCE to a collection of information unferse it contains a valid OMS control number.

INFORMATION DISCLOSURE	Ξ
STATEMENT BY APPLICANT	٢
(Not for submission under 37 CFR 1.99	,

Application Number		10567162		
Filing Date		2006-03-14		
First Named Inventor Katsu		shinge Marui		
Art Unit		1723		
Examiner Name Kim,		Sun U.		
Attorney Docket Number		10089-29		

	U.S.PATENTS			Remove						
Examiner Cit Initial* No		Patent Number	Kind Code ¹	Issue Date	Name of Patentee or Applicant		Pages,Columns,Lines where Relevant Passages or Releva Figures Appear			
	1	3928204		1975-12-23	Thomas					
	2	4080296		1978-03-21	Clark	-2-3-				
If you wisi	h to a	dd additional U.S. Pate	nt citatio	n information p	lease click the	Add button.	_	Add		_
			U.S.P	ATENT APPL	CATION PUB	LICATIONS		Remove		_
Examiner Initial*	Cite No	Publication Number	Kind Code ¹	Publication Date	Name of Pat of cited Doc	tentee or Applicant urnent	Releva	,Columns,L ant Passage s Appear		
If you wis	h to a	dd additional U.S. Publ		plication citation			d buttor	Add		_
Examiner Initial*	Cite No	Foreign Document Number ³	Country Code ²	Kind		Name of Patente Applicant of cited Document	e or	Pages,Colu where Rele Passages of Figures App	vant r Relevant	T6
	1	0873779	EP	A2	1998-10-28	Toyo Boseki Kabus Kaisha	shiki			
	2	2002292213	.IP		2002-10-08	KITZ CORP				Þ

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		10567162		
iling Date		2006-03-14		
irst Named Inventor	Katsu	shinge Marui		
Art Unit		1723		
Examiner Name Km, S		Sun U.		
Attorney Docket Number		10089-29		

If you wish	h to a	dd add	fitional Foreign Patent Document citation information please click the Add button	n Add	
			NON-PATENT LITERATURE DOCUMENTS	Remove	
Examiner Initials*	Examiner Cite Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (look, magazine, journal, sorial, symposium, catalog, etc), date, pages(s), volume-issue number(s), publisher, city and/or country where published.				ТБ
1 Supplementary Search Report issued March 19, 2007 in EP03817783.7-2113, PCT.IP0312194		14			
If you wish	h to a	dd add	fitional non-patent literature document citation information please click the Add I	outton Add	
			EXAMINER SIGNATURE		
Examiner	Signa	ture	Date Considered		
			reference considered, whether or not citation is in conformance with MPEP 609		

1 See Kind Codes of USPTO Patent Documents at www.ISPTO.000/c of MPE 90104. 2 Enter office that issued the document, by the hor-letter code (WIPO Standard ST3.) 2 For Jungsness petant counters, the includes not not be year of the insper or the protects the sent antimor of the patent of the paten

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		10567162		
Filing Date		2006-03-14		
First Named Inventor	Katsu	shinge Marui		
Art Unit		1723		
Examiner Name	Kim, \$	Sun U.		
Attorney Docket Number		10089-29		

CERTIFICATION STATEMENT

Please see 37	CFR 1.97 and	1.98 to make the a	ppropriate selection(s):
---------------	--------------	--------------------	--------------------------

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filling of the information disclosure statement. See 37 CFR 1.97(e/11).

OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquity, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 175(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 179(c).

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

Signature	/Cassandra T. Swain, Ph.D./	Date (YYYY-MM-DD)	2007-05-02
Name/Brint	Consender T. Comin. Dh. D.	Dogistration Number	49261

This collection of information is required by 3T CFR 1.87 and 1.98. The information is required to obtain or retain a benefit by the public which is to file fand by the USPTO to process) an application. Confidentially is governed by \$5 U.S. C. 12.9 and 3T CFR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Tradenary (Tifice, U.S. perpartment of Commence, P. 0. Box 1450, Alexandria, V.S. 2231-1450, D. NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA.2231-1450.

Privacy Act Statement

The Privacy Act of 1974 (P. L. 93-579) requires that you be given certain information in connection with your submission of the stackhold from related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, places be advised that (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) familishing of the information solicided is couldrain; and (3) the primoral pursuance for which the information is used by the U.S. Patient and Trademan Coffice is to process and/or examine your submission related to a patient agricultant or patient. If you do not furnish the requested process and/or examine your submission related to a patient agricultant or patient. If you do not furnish the requested results of the patient of the patient and the patient of the patient

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
 - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement negotiation.
 - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record pertains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
 - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974, as amended, pursuant to 5 U.S.C. 552(m).
 - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
 to the Patent Cooperation Treaty.
 - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
 - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uturing an insection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 4d U.S.C. 2904 and 2905. Such disclosure shall be made in accordance with the GSA requisions governing inseption of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the
 application pursuant to 35 U.S.C. 12(2) or issuance of a patent pursuant to 35 U.S.C. 151. Further, a record may be
 disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record was filled in application
 which became abandoned or in which the proceedings were terminated and which application is referenced by either a
 published application, an application open to public inspections or as issued patent.
 - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.